



INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification ⁴ : A61L 15/06	A1	(11) International Publication Number: WO 88/ 01878 (43) International Publication Date: 24 March 1988 (24.03.88)
(21) International Application Number: PCT/GB87/00658 (22) International Filing Date: 21 September 1987 (21.09.87) (31) Priority Application Numbers: 8622695 8629231 8701434 (32) Priority Dates: 20 September 1986 (20.09.86) 6 December 1986 (06.12.86) 22 January 1987 (22.01.87) (33) Priority Country: GB (71) Applicant (for all designated States except US): SMITH & NEPHEW ASSOCIATED COMPANIES PLC [GB/GB]; 2 Temple Place, Victoria Embankment, London WC2R 3BP (GB). (72) Inventor; and (75) Inventor/Applicant (for US only) : ANSELL, Christopher, Wilson, Guy [GB/GB]; 6 Woodland Road, Sawston, Cambridge CB2 4DU (GB).	(74) Agent: COLE, W., G.; Smith & Nephew Research Limited, Gilston Park, Harlow, Essex CM20 2RQ (GB). (81) Designated States: AT (European patent), AU, BE (European patent), CH (European patent), DE (European patent), FR (European patent), GB (European patent), IT (European patent), JP, LU (European patent), NL (European patent), SE (European patent), US. Published <i>With international search report. Before the expiration of the time limit for amending the claims and to be republished in the event of the receipt of amendments.</i>	
(54) Title: ADHESIVE DRESSING (57) Abstract Skin friendly adhesive dressings suitable for application such as wound dressings including first aid dressings, include a support layer which may be a knitted, woven or non-woven fabric or a microporous or continuous polymeric film, onto which is coated an inherently tacky gel or foam polyurethane adhesive. The adhesive contains less than 10 % by weight of materials leachable into water.		

FOR THE PURPOSES OF INFORMATION ONLY

Codes used to identify States party to the PCT on the front pages of pamphlets publishing international applications under the PCT.

AT	Austria	FR	France	ML	Mali
AU	Australia	GA	Gabon	MR	Mauritania
BB	Barbados	GB	United Kingdom	MW	Malawi
BE	Belgium	HU	Hungary	NL	Netherlands
BG	Bulgaria	IT	Italy	NO	Norway
BJ	Benin	JP	Japan	RO	Romania
BR	Brazil	KP	Democratic People's Republic of Korea	SD	Sudan
CF	Central African Republic	KR	Republic of Korea	SE	Sweden
CG	Congo	LI	Liechtenstein	SN	Senegal
CH	Switzerland	LK	Sri Lanka	SU	Soviet Union
CM	Cameroon	LU	Luxembourg	TD	Chad
DE	Germany, Federal Republic of	MC	Monaco	TG	Togo
DK	Denmark	MG	Madagascar	US	United States of America
FI	Finland				

- 1 -

ADHESIVE DRESSING

The present invention relates to adhesive dressings. More particularly the invention is for dressings utilizing skin-friendly inherently tacky polyurethane gel or foam adhesives such as would be required for wound care applications.

The use of polyurethane - based adhesives for skin application has been proposed, although not for applications where it is necessary for there to be substantially little or no leaching of materials from the adhesive and where the adhesive is required to be skin friendly.

SUBSTITUTE SHEET

- 2 -

In British Patent Nos 146817 and 1351909, there are described adhesive formulations, which may be inter alia based on polyurethanes which contain essentially leachable medicated extracts. Although such adhesives may be tacky, there is no suggestion that such formulations are gel or foam adhesives of inherent tackiness and a low level of leachable materials.

European Patent Specification No. 0147588 discloses self adhesive sheet-like structures comprising a support layer at least partially covered with a gel adhesive containing a mixture of a cross-linked polyurethane and immobilised polyhydroxy compounds having an average functionality if at least 3. However, such adhesives contain considerable amounts of leachable materials, in some cases as high as 50% and would not be suitable for use in wound areas of high exudation.

The present invention seeks to provide an adhesive dressing in which the adhesive is skin friendly i.e. causes no tendency to strip skin layers when removed, leaves no deposits, and yet can be employed for direct wound contact and contains substantially no leachable materials.

SUBSTITUTE SHEET

- 3 -

According to the present invention there is provided an adhesive dressing suitable for application to skin including a support layer for a coating of a polyurethane gel or foam adhesive, which adhesive is inherently tacky and contains not more than 10% by weight of leachable materials.

Tack is a property of an adhesive whereby light contact with the surface of another body brings about a condition requiring force to restore the original separated state, that is a property which will inhibit but not wholly prevent the removal of a smooth contacting surface such as a metal probe or a human finger. By inherent tack means the gel or foam possesses this property without requiring the addition of any further adhesion promoting component such as a tackifier. Tack and how it may be measured are discussed in Chapter 3 of Handbook of Pressure-Sensitive Adhesive Technology Editor D. Satas, Van Nostrand Reinhold Co. Inc. New York, 1982.

Suitably the inherently tacky gel is a pressure sensitive adhesive, that is the adherency of the gel is dependent upon the force used to apply it to a surface. Other types of adhesive which comprise inherently tacky gels of highly hydrophilic polyurethane and inherently

SUBSTITUTE SHEET

- 4 -

tacky foams of highly hydrophilic polyurethane are however, also within the scope of this invention.

The dressings of the invention are further characterised in that the content of materials leachable into water will be no greater than 10% by weight of the adhesive. More suitably the content of materials leachable into water will be less than 5%, preferably less than 2.5% by weight of the adhesive. The adhesives used in the dressings of the invention do not require the presence of additives such as humectants or plasticizers which may leach out into a wound surface and thus be considered generally unsatisfactory.

The adhesives employed in the dressings of the invention may be hydrophilic polyurethanes. One class of highly hydrophilic polyurethanes are those prepared by reacting water of a polyol with an isocyanate prepolymer which is the reaction product of a polyfunctional isocyanate and a polyoxyalkylene diol monoalkyl ether.

By highly hydrophilic is meant that the polyurethane when hydrated may contain from 35 to 95% by weight of water, aptly 50 to 92%, preferably 70 to 90% and more preferably 75 to 85% by weight water. Water absorption can be obtained by taking a known weight of

SUBSTITUTE SHEET

- 5 -

the polyurethane and immersing in water for 24 hours. The hydrated polymer is removed from the water, excess water is removed by lightly blotting with absorbent paper and then the hydrated polyurethane is weighed. The water absorption of the polyurethane (% by weight) can then be calculated as

$$\frac{(\text{weight of hydrated polyurethane} - \text{weight of dry polyurethane}) \times 100}{\text{weight of hydrated polyurethane}}$$

The inherently tacky adhesive gels or foams may be formed by mixing a prepolymer formed as the reaction product of polyisocyanate and polyoxyalkylene diol mono alkyl ether with a molar excess of water or with a stoichiometric amount of polyol or polyol mixture. The polyol is preferably a diol or mixture of diols. The excess of water used is absorbed by the tacky gel or foam produced.

Another class of adhesives suitable for use in the present invention are polyurethane adhesives which also contain acrylate residues in addition to the polyurethane residues. Such adhesives may be prepared by reacting an isocyanate prepolymer, which itself may be the reaction product of a polyfunctional isocyanate

SUBSTITUTE SHEET

- 6 -

and a polyoxy alkylene diol monoalkyl ether, with a hydroxy containing compound, e.g. water or a polyol and a hydroxyl - containing ester of acrylic or methacrylic acid and/or mixtures of such hydroxyl - containing compounds.

Suitable polyoxyalkylene diol mono alkyl ethers for use in forming the prepolymers employed to prepare the adhesives employed in the present invention include those in which the alkyl group contains from 1 to 18 carbon atoms and more suitably 2 to 6 carbon atoms and preferably 2 to 12 carbon atoms for example 4 carbon atoms that is the mono alkyl ether is a monobutyl ether.

Suitably the polyoxyalkylene residue in the mono alkyl ether is a hydrophilic residue that contains polyoxyethylene or polyoxypropylene residues or mixtures thereof. Preferred polyoxyalkylene residues are those which contain a mixture of polyoxyethylene and polyoxypropylene residues in a ratio of from 20:80 to 80:20 for example 50:50. The residues may be randomly arranged with respect to each other.

A preferred polyoxyalkylene diol mono alkyl ether is therefore polyoxyethylene - polyoxypropylene mono

- 7 -

butyl ether in which the ratio of polyoxyethylene to polyoxypropylene residues is about 50:50 such as the Emkarox polyols sold by I.C.I. plc.

Aptly the molecular weight of the monoalkyl ether is from 3000 to 12000, and more suitably is 4000 to 10000 and preferably is 5000 to 9000. It has been found that polyoxyalkylene diol monoalkyl ethers within this molecular weight range provide polymers which are tacky and suitable for use on the skin while use of lower molecular weight materials tend to provide polymers which are not suitable for use as adhesives because they are insufficiently tacky.

The polyoxyalkylene diol mono alkyl ethers for use in forming the prepolymer will normally contain water. It is preferred however that the polyoxyalkylene diol monoalkyl ether contains less than 1% by weight of water so as to limit the proportion of urea groups formed in the reaction with the polyisocyanate.

The polyisocyanate used for forming the prepolymer will have a functionality of greater than 2 and may suitably have a functionality of from 2.1 to 5 and more suitably from 2.2 to 3.5 and preferably from 2.5 to 3.0 for example 2.5, 2.7, 2.85 or 2.9. Suitable

- 8 -

polyisocyanates include (cyclo) aliphatic and aromatic polyisocyanates. Preferred polyisocyanates are aromatic polyisocyanates for example those based on a polymeric methylene di phenyl diisocyanate, for example the Suprasecs (trade mark), which are available from I.C.I. It is preferred that the functionality of the isocyanate is not more than 3 as this leads to a higher cross-link density and is manifest ultimately as a harder adhesive which may not be advantageous for an adhesive which is to be applied to the skin.

The prepolymers may be simply prepared by heating the two components together in the required proportions at an elevated temperature for sufficient time for the reaction to be completed, for example 90°C for 2 hours, in the presence of 0.2% w/w catalyst of dibutyl tin dilaurate (T12). The mole ratio NCO/OH is suitably from 2.0 to 4.0 in this reaction.

The adhesives based solely on polyurethanes may be prepared by reacting the prepolymer described above with water. The reaction mixture may contain from 10% w/w water and 90% w/w prepolymer to 90% w/w water and 10% w/w prepolymer. It is preferred, however, when forming a tacky gel that the concentration of prepolymer is less than 25% w/w. If the concentration of prepolymer is

SUBSTITUTE SHEET

- 9 -

greater than 25% w/w then a tacky foam is produced, which also forms part of this invention. Normally the reactants are mixed together at room temperature and allowed to cure for at least 30 minutes. The excess water is taken up into the tacky gel or foam to produce a swollen gel or foam. It has been found that these gels and foams may take up a further amount of water often upto a weight equal to their original weight.

The polyurethane adhesives employed in the present invention may also be prepared by reacting the prepolymer with a stoichiometric amount of polyol. The reactants are mixed together and cured at elevated temperature for 15 minutes to 2 hours. The resultant polyurethane polymer is in the form of a dry adhesive mass which is inherently tacky and may absorb upto 90% by weight of water. Suitably the polyol is a diol. The choice of the polyol and isocyanate-content of the prepolymer govern the degree of tackiness of the resulting adhesive. Thus the lower isocyanate-content of the prepolymer, the lower is the cross-link density of the resultant polymer and the more tacky is the adhesive. Thus, also the higher the molecular weight of the polyol the softer is the adhesive mass.

SUBSTITUTE SHEET

- 10 -

Suitable polyols are diols which include alkylene glycols, di-alkylene glycols and polyoxyalkylene diol block copolymers. Suitable diols include diethylene glycol, polyoxyethylene diol having a molecular weight of about 200 and a polyoxyethylene-polyoxypropylene diol block copolymer having a molecular weight of about 2600, and polyoxypropylene diol of molecular wt about 2000.

The polyurethane-acrylate adhesives employed in the present invention may be prepared by reacting the prepolymer described above with hydroxy-containing compounds a proportion of which also contains an unsaturated functionality which is reactive when exposed to irradiation for example ultra violet or electron beam irradiation.

Suitable unsaturated compounds include esters of acrylic or methacrylic acid in which there is at least one hydroxyl functional group which is capable of reacting with isocyanate present. Preferred esters include hydroxy ethyl methacrylate and methacrylate mono-esters of polyoxylalkylene diols in which the number of repeating ether units is from 1 to 25 and preferably 2 to 10 for example 6.

SUBSTITUTE SHEET

- 11 -

The remainder of the hydroxyl-containing compounds comprise polyols such as diols or triols or monools such as primary alcohols or polyoxyalkylene monoalkyl ethers as herein before described. Preferably the hydroxyl-containing compounds are polyoxyalkylene monoalkyl ethers or other mono-ols which may confer additional tacky properties is the adhesive. Such mono-ols include hydrogenated mono hydroxylin tackifying resins such as hydrogenated abietyl alcohol.

The proportion of hydroxyl-containing compound which also contains an unsaturated functionality is such as to react with from 15 to 25% of the free isocyanate groups in the prepolymer. The remainder of the free isocyanate groups react with the other hydroxyl-containing compounds.

The resulting polymer has therefore pendant unsaturated groups which are capable of interacting with each other to cross-link the polymer under the influence of a polymerisation initiator and irradiation. One suitable form of radiation is ultraviolet irradiation. The polymer may be mixed with a photoinitiator and the mixture irradiated by means of ultraviolet radiation. This causes the polymer to become cross-linked.

SUBSTITUTE SHEET

- 12 -

The polyurethane polymer so formed is a cross-linked polymer which is capable of absorbing from 5 to 95% by weight of water depending upon the reactants employed.

Alternatively the adhesive may be formed as a 'wet' adhesive by dispersing the uncross-linked polymer and photoinitiator in an appropriate volume of water and then irradiating to form the adhesive. Suitably the wet adhesive may contain from 40 to 65% by weight of water.

The backing or supporting layers employed in the present invention should be capable of conforming to the body contours when applied to the skin and should be flexible enough to move with the body without becoming detached. Suitable flexible backing materials include knitted, woven or non-woven fabrics, nets, microporous films such as plasticised polyvinyl chloride films, polymer blend films containing voids, polymeric films, including thermoplastic polyurethane and hydrophilic polyurethane, elastomeric polyesters, styrene-butadiene block copolymers such as Kraton (Trade mark) thermoplastic rubbers.

Favoured materials which may be used as the backing layer in the dressing embodiment of the present invention include films of hydrophilic polymers. Apt

SUBSTITUTE SHEET

- 13 -

hydrophilic polymers include hydrophilic polyurethanes, polyvinyl pyrrolidone, polyvinyl alcohol and cellulosic derivatives.

A favoured hydrophilic polymer is a hydrophilic polyurethane. Suitable hydrophilic polyurethanes include those having the composition and prepared by the process described in British Patent No. 2093190B. The most suitable hydrophilic polyurethanes are those which contain from 5 to 50% by weight of water when hydrated move suitably to 10 to 40% by weight of water and which have a thickness when present in a dressing of from 25 to 120 μ m, more suitably 30 to 60 μ m. A preferred film of hydrophilic polyurethane has a water content when hydrated of 20 to 30% for example 25% and a thickness of 50 μ m for example 30 μ m.

A favoured flexible backing material is a microporous plasticised polyvinyl chloride film formed by the process disclosed in British Patent No. 884232. The preferred microporous plasticised polyvinyl chloride 30 to films have a thickness of from 100 to 300 μ m, for example 150 μ m, 200 μ m and 250 μ m.

A further favoured backing layer is a thermoplastic polyurethane including the linear polyester

SUBSTITUTE SHEET

- 14 -

polyurethanes or polyether polyurethanes known as Estanes (Trade mark). Such polyurethanes are used as films from 15 to 75 μm in thickness, more favourably 20 to 35 μm in thickness for example about 25 μm or 30 μm .

The adhesive layer may have a thickness of from 0.05 to 5mm depending upon the type of skin lesion, wound or other similar use the dressing is to be put, that is thinner layers may be used on non-exuding wounds or on dressings for use at intravenous sites while thicker layers may be used on exuding wounds where the absorptive properties of the adhesive may be advantageously used. More suitably the adhesive layer may have a thickness of 1.5 to 4mm and preferably 2 to 3mm.

The dressings of the present invention may be prepared by casting the backing layer from a solution of the appropriate polymer at a suitable concentration onto a silicone release paper and removing the solvent to give a film of the required thickness. A film of the adhesive at the required thickness is prepared and the two films brought together and adhered to each other to form a laminate, preferably without the use of any further adhesives. A release coated protector may be placed over the adhesive layer and the laminate cut into

SUBSTITUTE SHEET

- 15 -

dressings of the appropriate size for example 5cm x 5cm, 7.5cm x 7.5cm, 8cm x 8cm, 10cm x 10cm, 5cm x 10cm and 10cm x 20cm. The dressings may be packaged in a bacteria-proof and water-proof pouch and be sterilized by conventional methods such as irradiation and ethylene oxide.

In use the sterile dressing is removed from the pouch, the protector removed from the adhesive surface of the dressing and the dressing is applied to the wound.

The dressing may be used as a dressing for wounds caused by physical or surgical trauma, burns, ulcers and the like, as a surgical drape, as a dressing for intravenous access sites and any dressing for which long term attachment to the skin may be required.

In a further aspect therefore the present invention provides a method of treating a wound on an animal body by applying to the wound an adhesive dressing as hereinbefore described.

Many medicinal agents may be incorporated into the adhesives of the present invention. By medicinal agent it is meant pharmacologically active agents and agents

SUBSTITUTE SHEET

- 16 -

including topical anaesthetics such as xylocaine, bacteriostatic agents such as silver nitrate; antibacterial agents of which preferred agents are silver sulphadiazine and chlorhexidine salts; antibiotics; topical steroids, enzymes, tissue stimulants, coagulants and anticoagulants and antifungal agents. Other agents such as emollients may be added after the reaction step.

Advantageously water soluble medicaments such as chlorhexidine and its salts may be dissolved in the water which is used to react with the prepolymer. It is found that chlorhexidine is unaffected during the process and the resulting adhesive provides effective release of chlorhexidine when placed onto the skin.

Example 1

A polyoxyethylene - polyoxypropylene diol monobutyl ether, which has a ratio of polyoxyethylene to polyoxypropylene residues of 1:1 and molecular weight of 4095 (300g, 0.073 moles based on OH value) and a polymeric methylene diphenyl diisocyanate (37.21g, 0.266 moles, - NCO functionality of 2.7) were mixed together at an NCO/OH ratio of 2.5 in a 700cm³ flange flask fitted with an overhead stirrer. The flask was heated

- 17 -

in a water bath set to a temperature of 90°C. A catalyst comprising dibutyl tin dilaurate (0.2% w/w) was added. The mixture was stirred at 90°C for two hours. The prepolymer so formed was allowed to cool. The prepolymer was a golden yellow viscous liquid which may be stored in a capped bottle until ready for use. The prepolymer was found to have an isocyanate content of 1.98%.

A portion of the prepolymer and the calculated quantity of diethylene glycol which would react with all the available isocyanate were mixed at room temperature until homogeneous and then spread onto a silicone release paper at a weight per unit area of 280gsm and cured at 90°C to give an adhesive mass.

The adhesive mass contained 85% by weight of water when fully hydrated.

A precast film of a thermoplastic polyether polyurethane may be transfer coated onto the adhesive mass and the laminate strip so formed cut into pieces which are suitable for adhesive dressings and packed in a bacteria proof and waterproof package and sterilised by irradiation.

SUBSTITUTE SHEET

- 18 -

In use the sterile dressing is removed from the package, the silicone coated release paper removed and the adhesive layer placed against and around the wound area.

A film of adhesive gel was coated onto various substrates commonly used in wound dressings at either a weight per unit area of either 285 gsm or 70 gsm. The upright and inverted moisture transmission rates were measured and the results shown in the following table.

Demonstration 1

Upright MVP Demonstration

Discs of material under test are clamped over Payne Permeability Cups (flanged metal cups) using sealing rings and screw clamps. The exposed surface area of the test sample may be conveniently 10cm². Each cup contains approximately 10ml of distilled water.

After weighing the cups are placed in a fan assisted electric oven which is maintained at $37 \pm 1^{\circ}\text{C}$. The relative humidity within the oven is maintained at

SUBSTITUTE SHEET

- 19 -

approximately 10% by placing 1Kg of anhydrous 3-8 mesh calcium chloride on the floor of the oven.

The cups are removed after 24 hours, allowed to cool for 20 minutes and re-weighed. The MVP of the test material is calculated from the weight loss experienced in units of grams of weight per square metre per 24 hours.

Inverted MVP Determination

The method described above is employed except that the Payne Cups are inverted in the oven so that the water within the cups is in contact with the test material and in this case with the gel adhesive.

SUBSTITUTE SHEET

- 20 -

Substrate	Weight per unit area of adhesive	Upright (U)/ Inverted cup (I)	MVTR (g/m ² /2h)
Polyurethane containing 25% by weight water when hydrated, 30gsm	285	U	1472
	285	I	7356
Polyurethane (as above)	70	U	1455
	70	I	>14000
Plasticised Polyvinyl Chloride (150µm thick)	70	U	1444
	70	I	5384
Knitted Fabric	70	U	5600
	70	I	>14000
Polymer net prepared according to UK Patent No. 1531715	70	U	1644
	70	I	>14000

SUBSTITUTE SHEET

- 21 -

CLAIMS

1. An adhesive dressing suitable for application to skin including a support layer for a coating of a polyurethane gel or foam adhesive, which adhesive is inherently tacky and contains not more than 10% by weight of leachable materials.
2. A dressing as claimed in claim 1 wherein the polyurethane is a hydrophilic polyurethane.
3. A dressing as claimed in claim 2 wherein, when hydrated, the polyurethane contains 35 to 95% by weight water.
4. A dressing as claimed in claim 3 wherein, when hydrated the polyurethane contains from 75 to 85% by weight water.
5. A dressing as claimed in any one of the peeling claims wherein the adhesive contains residues derived from both polyurethane and an acrylate or methacrylate.
6. A dressing as claimed in any one of the preceeding claims wherein the polyurethane residues are derived from an isocyanate prepolymer which is the reaction

SUBSTITUTE SHEET

- 22 -

product of polyisocyanate and a polyoxyalkylene diol monoalkyl ether.

7. A dressing as claimed in claim 6 wherein the hydroxy compound is water or a polyol.

9. A dressing as claimed in claim 7 wherein at least a proportion of the hydroxy compound hydroxyl compound which is capable of reacting with isocyanate and which contains an unsaturated functionality which is reactive when exposed to irradiation.

10. A dressing as claimed in claim 9 wherein the hydroxyl compound in an ester of acrylic or methacrylic acid.

11. A dressing as claimed in any one of claims 8 to 10 wherein the concentration of water is from 10 to 90% by weight of the prepolymer.

12. A dressing as claimed in claim 11 wherein the water is less than 25% by weight of the prepolymer.

13. A dressing as claimed in any one of claims 8 to 12 wherein the polyol is a diol.

SUBSTITUTE SHEET

- 23 -

14. A dressing as claimed in claim 13 wherein the diol is present in a stoichiometric amount with respect to the isocyanate prepolymer.

15. A dressing as claimed in any one of claims 6 to 14 wherein the alkyl moiety of the polyoxyalkylene monoalkyl ether contain from 2 to 6 carbon atoms.

16. A dressing as claimed in any one of claims 6 to 15 wherein the polyoxyalkylene residue of the ether in a hydrophilic residue containing polyoxyethylene or polyoxypropylene residues or mixtures thereof.

17. A dressing as claimed in either of claims 15 or 16 wherein the ether is a polyoxyethylene - polyoxypropylene mono-butyl ether in which the ratio of polyoxyethylene to polyoxypropylene residues is about 50:50.

18. A dressing as claimed in any one of the preceding claims wherein the support layer is a flexible knitted woven or non-woven fabric, a net, a microporous polymer film, or a moisture vapour permeable polymeric film.

19. A dressing as claimed in any one of the preceding claims wherein the adhesive contains a pharmacologically active medicinal agent.

SUBSTITUTE SHEET

- 24 -

20. A dressing as claimed in any one of the preceeding claims wherein the content of materials leachable into water is less than 2.5% by weight of the adhesive.

21. A dressing as claimed in any one of the preceeding claims, within a sterile package.

22. A method of treating a wound on an animal body by applying to the wound an adhesive dressing as claimed in any one of the preceeding claims.

SUBSTITUTE SHEET

INTERNATIONAL SEARCH REPORT

International Application No PCT/GB 87/00658

I. CLASSIFICATION OF SUBJECT MATTER (if several classification symbols apply, indicate all) ⁴ According to International Patent Classification (IPC) or to both National Classification and IPC IPC ⁴ : A 61 L 15/06											
II. FIELDS SEARCHED <div style="text-align: right; margin-right: 100px;">Minimum Documentation Searched ⁷</div> <table style="width: 100%; border: none;"> <tr> <td style="width: 30%; border-bottom: 1px solid black;">Classification System</td> <td style="border-bottom: 1px solid black;">Classification Symbols</td> </tr> <tr> <td style="border: none;">IPC⁴</td> <td style="border: none;">A 61 L 15/06</td> </tr> </table> <div style="text-align: center; margin-top: 10px;"> Documentation Searched other than Minimum Documentation to the extent that such Documents are included in the Fields Searched ⁸ </div>			Classification System	Classification Symbols	IPC ⁴	A 61 L 15/06					
Classification System	Classification Symbols										
IPC ⁴	A 61 L 15/06										
III. DOCUMENTS CONSIDERED TO BE RELEVANT ⁹ <table border="1" style="width: 100%; border-collapse: collapse;"> <thead> <tr> <th style="width: 10%;">Category ⁹</th> <th style="width: 70%;">Citation of Document, ¹¹ with indication, where appropriate, of the relevant passages ¹²</th> <th style="width: 20%;">Relevant to Claim No. ¹³</th> </tr> </thead> <tbody> <tr> <td style="text-align: center; vertical-align: top;">A</td> <td>EP, A, 0147588 (BAYER) 10 July 1985 cited in the application --</td> <td></td> </tr> <tr> <td style="text-align: center; vertical-align: top;">A</td> <td>EP, A, 0080008 (CARL FREUDENBERG) 1 June 1983 -----</td> <td></td> </tr> </tbody> </table>			Category ⁹	Citation of Document, ¹¹ with indication, where appropriate, of the relevant passages ¹²	Relevant to Claim No. ¹³	A	EP, A, 0147588 (BAYER) 10 July 1985 cited in the application --		A	EP, A, 0080008 (CARL FREUDENBERG) 1 June 1983 -----	
Category ⁹	Citation of Document, ¹¹ with indication, where appropriate, of the relevant passages ¹²	Relevant to Claim No. ¹³									
A	EP, A, 0147588 (BAYER) 10 July 1985 cited in the application --										
A	EP, A, 0080008 (CARL FREUDENBERG) 1 June 1983 -----										
<div style="display: flex; justify-content: space-between;"> <div style="width: 45%;"> <p>¹⁰ Special categories of cited documents:</p> <p>"A" document defining the general state of the art which is not considered to be of particular relevance</p> <p>"E" earlier document but published on or after the international filing date</p> <p>"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)</p> <p>"O" document referring to an oral disclosure, use, exhibition or other means</p> <p>"P" document published prior to the international filing date but later than the priority date claimed</p> </div> <div style="width: 45%;"> <p>"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention</p> <p>"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step</p> <p>"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.</p> <p>"Δ" document member of the same patent family</p> </div> </div>											
IV. CERTIFICATION <table style="width: 100%; border: none;"> <tr> <td style="width: 50%; border-bottom: 1px solid black;">Date of the Actual Completion of the International Search</td> <td style="width: 50%; border-bottom: 1px solid black;">Date of Mailing of this International Search Report</td> </tr> <tr> <td style="border: none;">24th November 1987</td> <td style="border: none; text-align: center;">14 JAN 1988</td> </tr> <tr> <td style="border-bottom: 1px solid black;">International Searching Authority</td> <td style="border-bottom: 1px solid black;">Signature of Authorized Officer</td> </tr> <tr> <td style="border: none; text-align: center;">EUROPEAN PATENT OFFICE</td> <td style="border: none; text-align: center;"> P.C.G. VAN DER PUTTEN </td> </tr> </table>			Date of the Actual Completion of the International Search	Date of Mailing of this International Search Report	24th November 1987	14 JAN 1988	International Searching Authority	Signature of Authorized Officer	EUROPEAN PATENT OFFICE	 P.C.G. VAN DER PUTTEN	
Date of the Actual Completion of the International Search	Date of Mailing of this International Search Report										
24th November 1987	14 JAN 1988										
International Searching Authority	Signature of Authorized Officer										
EUROPEAN PATENT OFFICE	 P.C.G. VAN DER PUTTEN										

**ANNEX TO THE INTERNATIONAL SEARCH REPORT
ON INTERNATIONAL PATENT APPLICATION NO.**

GB 8700658

SA 18772

This annex lists the patent family members relating to the patent documents cited in the above-mentioned international search report. The members are as contained in the European Patent Office EDP file on 22/12/87
The European Patent Office is in no way liable for these particulars which are merely given for the purpose of information.

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
EP-A- 0147588	10-07-85	AU-A- 3546684	23-05-85
		DE-A- 3341555	30-05-85
		JP-A- 60118783	26-06-85
		US-A- 4661099	28-04-87
EP-A- 0080008	01-06-83	JP-A- 58089261	27-05-83